abcam

Product datasheet

Recombinant Human Factor X protein ab158407

1 Image

Description

Product name Recombinant Human Factor X protein

Expression system Wheat germ

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Sequence NNILARVTRANSFLEEMKKGHLERECMEETCSYEEAREVF

EDSDKTNEFW

NKYKDGDQCETSPCQNQGKCKDGLGEYTCTCLEGFEGK

NCELFTRKLCSL

DNGDCDQFCHEEQNSVVCSCARGYTLADNGKACIPTGPY

PCGKQTLERRK

RSVAQATSSSGEAPDSITWKPYDAADLDPTENPFDLLDF

NQTQPERGDNN

LTRIVGGQECKDGECPWQALLINEENEGFCGGTILSEFYIL

TAAHCLYQA

KRFKVRVGDRNTEQEEGGEAVHEVEVVIKHNRFTKETYD

FDIAVLRLKTP

ITFRMNVAPACLPERDWAESTLMTQKTGIVSGFGRTHEKG

RQSTRLKMLE

VPYVDRNSCKLSSSFIITQNMFCAGYDTKQEDACQGDSG

GPHVTRFKDTY

FVTGIVSWGEGCARKGKYGIYTKVTAFLKWIDRSMKTRGL

PKAKSHAPEV ITSSPLK

Amino acids 32 to 488

Tags GST tag N-Terminus

Specifications

Our <u>Abpromise guarantee</u> covers the use of ab158407 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications Western blot

ELISA

1

Form

Liquid

Additional notes

Preparation and Storage

Stability and Storage

Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.

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Constituents: 0.31% Glutathione, 0.79% Tris HCI

General Info

Function Factor Xa is a vitamin K-dependent glycoprotein that converts prothrombin to thrombin in the

presence of factor Va, calcium and phospholipid during blood clotting.

Tissue specificity

Plasma; synthesized in the liver.

Involvement in disease Defects in F10 are the cause of factor X deficiency (FA10D) [MIM:227600]. A hemorrhagic

disease with variable presentation. Affected individuals can manifest prolonged nasal and

mucosal hemorrhage, menorrhagia, hematuria, and occasionally hemarthrosis. Some patients do

not have clinical bleeding diathesis.

Sequence similaritiesBelongs to the peptidase S1 family.

Contains 2 EGF-like domains.

Contains 1 Gla (gamma-carboxy-glutamate) domain.

Contains 1 peptidase S1 domain.

Post-translational

modifications

The vitamin K-dependent, enzymatic carboxylation of some glutamate residues allows the

modified protein to bind calcium.

N- and O-glycosylated.

The activation peptide is cleaved by factor IXa (in the intrinsic pathway), or by factor VIIa (in the

extrinsic pathway).

The iron and 2-oxoglutarate dependent 3-hydroxylation of aspartate and asparagine is (R)

stereospecific within EGF domains.

Cellular localization

Secreted.

Images

175 – 83 – 62 – 47.5 – 32.5 – 25 – 25 – SDS-PAGE - Recombinant Human Factor X protein (ab158407)

ab158407 on a 12.5% SDS-PAGE stained with Coomassie Blue.

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